US ERA ARCHIVE DOCUMENT

## UNITED STATES ENVIRONMENTAL PROTECTION AGENCY



WASHINGTON, D.C. 20460

OFFICE OF PREVENTION, PESTICIDES AND TOXIC SUBSTANCES

SPINS 2/15/00

% by wt.

January 21, 2000

## MEMORANDUM:

Subject:

EPA Reg. No.: 10182-334/Talon-G Rodenticide Bait Pack (Mini-Pellets)

EPA Reg. No.: 10182-335/Talon-G Rodenticide Bait Pack Pellets w/Bitrex

EPA Reg. No.: 10182-336/Brodifacoum Rodenticide Pellets w/Bitrex

EPA Reg. No.: 10182-337/HAVOC Rodenticide Bait Pack Pellets w/Bitrex EPA Reg. No.: 10182-338/HAVOC Rodenticide Bait Pack Mini-Pellets EPA Reg. No.: 10182-341/Talon-G Rodenticide Mini-Pellets w/Bitrex

Case No.: 2760

From:

Marianne Lewis, Biologist

Product Reregistration Branch

Special Review and Reregistration Division (7508C)

To:

Venus Eagle-Kunst, CRM

Product Reregistration Branch

Special Review and Reregistration Division (7508C)

Applicant:

ZENECA Inc.

1800 Concord Pike

Wilmington, DE 19897

FORMULATION FROM EPA Reg. No. 10182-334 LABEL:

Active Ingredient(s):
Brodifacoum: 3-[3-(4'-bromo-[1,1'-biphenyl]-4-yl)-1,2,3,4-tetrahydro1-napthalenyl]-4-hydroxy-2H-1-benzopyran-2-one 0.005%
Inert Ingredient(s): 99,995%
Total 100.000%

FORMULATION FROM EPA Reg. No. 10182-335 LABEL:	% by wt.
Active Ingredient(s):  Brodifacoum: 3-[3-(4'-bromo-[1,1'-biphenyl]-4-yl)-1,2,3,4-tetrahydro- 1-napthalenyl]-4-hydroxy-2H-1-benzopyran-2-one Inert Ingredient(s):  Total	0.005 % <u>99.995 %</u> 100.000 %
FORMULATION FROM EPA Reg. No. 10182-336 LABEL:	% by wt.
Active Ingredient(s):  Brodifacoum: 3-[3-(4'-bromo-[1,1'-biphenyl]-4-yl)-1,2,3,4-tetrahydro- 1-napthalenyl]-4-hydroxy-2H-1-benzopyran-2-one Inert Ingredient(s):  Total	0.005 % <u>99.995 %</u> 100.000 %
FORMULATION FROM EPA Reg. No. 10182-337 LABEL:	% by wt.
Active Ingredient(s): Brodifacoum: 3-[3-(4'-bromo-[1,1'-biphenyl]-4-yl)-1,2,3,4-tetrahydro- 1-napthalenyl]-4-hydroxy-2H-1-benzopyran-2-one Inert Ingredient(s):  Total	. 0.005 % . <u>99,995 %</u> 100.000 %
FORMULATION FROM EPA Reg. No. 10182-338 LABEL:	% by wt.
Active Ingredient(s): Brodifacoum: 3-[3-(4'-bromo-[1,1'-biphenyl]-4-yl)-1,2,3,4-tetrahydro- 1-napthalenyl]-4-hydroxy-2H-1-benzopyran-2-one Inert Ingredient(s):  Total	0.00570
FORMULATION FROM EPA Reg. No. 10182-341 LABEL:	% by wt.
Active Ingredient(s):  Brodifacoum: 3-[3-(4'-bromo-[1,1'-biphenyl]-4-yl)-1,2,3,4-tetrahydro- 1-napthalenyl]-4-hydroxy-2H-1-benzopyran-2-one Inert Ingredient(s):	0.003 //

BACKGROUND: In the 8 month response to the Rodenticide Cluster RED, the registrant is requesting to waive the acute inhalation study (81-3) for the following products: EPA Reg. Nos.: 10182-334, 10182-335, 10182-336, 10182-337, 10182-338, and 10182-341. The registrant conducted a trial run to "generate the test material, using the pelleted formulation, which met the minimum particle size distribution requirement in order to conduct an acute inhalation study according to EPA test protocol guidelines." The laboratory was unable to generate the appropriate percentage of particles in the acceptable size range of  $1-4\mu m$ . The registrant was contacted and an inquiry was made as to how long the test material was milled/ground. Based on the registrant's reply, it appears that the milling was conducted over the course of one business day, or approximately 8 hours. The Agency prefers that any test material that does not easily produce the appropriately sized particles, be milled or ground (by air mill, ball mill, hammer mill, utilization of dry ice) for 24 hours if necessary. If the laboratory can demonstrate that the particle size has reached a plateau prior to 24 hours, then no further milling is needed.

Since the laboratory only attempted to mill the test material for approximately 8 hours, the Agency can not conclude that respirable particles could not be generated from this product. However, based on information available to the Agency, the acute inhalation study will be waived for the subject products and all will be classified as Toxicity Category I.

## **RECOMMENDATIONS:**

• The acute inhalation study (81-3) requirement for the subject products has been waived. All subject products will be classified as Toxicity Category I.

The acute toxicity profile for EPA Reg. No. 10182-334 is currently:

Acute Oral Data Needed
Acute Dermal Data Needed
Acute Inhalation I Waived
Primary Eye Data Needed
Primary Dermal Data Needed
Skin Sensitization Data Needed

The acute toxicity profile for EPA Reg. No. 10182-335 is currently:

Acute Oral Data Needed
Acute Dermal Data Needed
Acute Inhalation I Waived
Primary Eye Data Needed
Primary Dermal Data Needed
Skin Sensitization Data Needed

The acute toxicity profile for EPA Reg. No. 10182-336 is currently:

Acute Oral Data Needed
Acute Dermal Data Needed
Acute Inhalation I Waived
Primary Eye Data Needed
Primary Dermal Data Needed
Skin Sensitization Data Needed

The acute toxicity profile for EPA Reg. No. 10182-337 is currently:

Acute Oral Data Needed
Acute Dermal Data Needed
Acute Inhalation I Waived
Primary Eye Data Needed
Primary Dermal Data Needed
Skin Sensitization Data Needed

The acute toxicity profile for EPA Reg. No. 10182-338 is currently:

Acute Oral Data Needed
Acute Dermal Data Needed
Acute Inhalation I Waived
Primary Eye Data Needed
Primary Dermal Data Needed
Skin Sensitization Data Needed

The acute toxicity profile for EPA Reg. No. 10182-341 is currently:

Acute Oral Data Needed
Acute Dermal Data Needed
Acute Inhalation I Waived
Primary Eye Data Needed
Primary Dermal Data Needed
Skin Sensitization Data Needed

NOTE: The labeling will be completed when all of the required information is received.